


 Cite this: *RSC Adv.*, 2018, **8**, 33065

Controlled hydrothermal temperature provides tunable permittivity and an improved electromagnetic absorption performance of reduced graphene oxide[†]

Yilu Xia,^{ab} Jiankun Wang,^c Chaochan Chen,^d Da Huo,^a Yue Wen,^a Wenyue Wang,^a Mengxiao Sun,^a Chang Xu,^e Aming Xie,^{ID}^a Fan Wu^{ID}^{*a} and Zhangqi Feng^{ID}^{*b}

Received 9th July 2018
 Accepted 24th August 2018

DOI: 10.1039/c8ra05843a
rsc.li/rsc-advances

Reduced graphene oxide (RGO) has been prepared by a hydrothermal reduction method to explore the effects of reaction temperature on its permittivity and electromagnetic absorption (EA) performance. This study shows that by controlling the oxygen functional groups on the RGO surface it is also possible to obtain an ideal EA performance without any other decorated nanomaterials.

Electromagnetic fields play a critical role in modern electronic and communications industries. However, they also cause humans and wildlife to become chronically exposed to excessive electromagnetic radiation in the environment. Electromagnetic absorption (EA) is one of the most effective methods of managing electromagnetic pollution. Due to the mechanism, EA performance mainly depends on the permittivity and permeability of a material.¹ Thus, dielectric and magnetic materials and their hybrids have been considered as fillers to fabricate polymer based composites with EA performance properties.^{2,3} Due to their large aspect ratios and fascinating dielectric properties, two-dimensional (2D) nanomaterials show huge potential for the application of EA.^{1,4-8} Among these 2D nanomaterials, the permittivity and EA performance of reduced graphene oxide (RGO) have been most widely studied. Well-reduced RGO can significantly enhance the permittivity of polymers,^{1,4,6} however, it easily causes impedance mismatch between the air and the composites.⁹ This phenomenon not only limits the EA performance of the composites, but also

makes the incident electromagnetic waves more likely to reflect off the surface of the composite. In order to remedy this defect, multi-component composites have been considered to decorate RGO to achieve ideal EA performances, such as ZnO,¹⁰ MoS₂,¹¹ NiFe₂O₄,¹² MoS₂@Fe₃O₄,¹³ CoNi@SiO₂,¹⁴ and V₂O₅/carbon nanotubes (CNTs).¹⁵

So far, only a few studies have focused on the regulation of permittivity and the improvement of EA performance of pure RGO.¹ However, the relationship between synthetic parameters and electromagnetic properties is worthy of being explored further. In this study, we discuss the effects of hydrothermal temperature on the permittivity and EA performance of RGO. The results show that regulation of the C/O ratio through controlling the hydrothermal temperature can provide tunable permittivity and an ideal EA performance.

Graphene oxide (GO) powder (70 mg) was evenly dispersed in distilled water (1 mg mL⁻¹) by ultrasonication. The dispersion was sealed in a 100 mL Teflon-lined autoclave and maintained at 100 °C for 6 hours to remove the oxygen functional groups, then naturally cooled to room temperature. The resulting precipitate from the reaction was then centrifuged. Then the platelets were collected and freeze-dried for 24 hours. The hydrothermal temperatures were 100, 120, 140, 160 and 180 °C, and we thus designated the RGO products as RGO₁₀₀, RGO₁₂₀, RGO₁₄₀, RGO₁₆₀ and RGO₁₈₀.

The micro-morphologies of the as-prepared RGO samples were investigated by SEM analysis and the images show that they have similar 2D flexible layer-like structures (Fig. S1†). However, with the increase in temperature, the RGO platelets formed large layers with irregular holes.

In the XRD spectra (Fig. 1a), the diffraction peaks of graphite (002) move gradually to 24° with the increase of hydrothermal temperature. A peak around 12.5° can be found in RGO₁₀₀, and it suggests that the partial GO has not been well-reduced. On the

^aSchool of Mechanical Engineering, Nanjing University of Science & Technology, Nanjing 210094, P. R. China. E-mail: wufan@njust.edu.cn

^bSchool of Chemical Engineering, Nanjing University of Science & Technology, Nanjing 210094, P. R. China. E-mail: fengzhangqi1981@163.com

^cPudong Foreign Language School Affiliated Shanghai International Studies University, Shanghai 201203, P. R. China

^dDivision of Electron and Electricity Measurement Technology, Shanghai Institute of Measurement and Testing Technology, Shanghai 201203, P. R. China

^eState Key Laboratory for Disaster Prevention & Mitigation of Explosion & Impact, Army Engineering University of PLA, Nanjing 210007, P. R. China

† Electronic supplementary information (ESI) available: Experimental section (S1); SEM images of RGO₁₀₀, RGO₁₂₀, RGO₁₄₀, RGO₁₆₀ and RGO₁₈₀ (Fig. S1); the atom% of C/O in RGO₁₀₀, RGO₁₂₀, RGO₁₄₀, RGO₁₆₀ and RGO₁₈₀ (Table S1); typical thickness dependent EA performance; RGO₁₄₀ and RGO₁₆₀ are loaded at 5 wt% and 7 wt% in wax composites (Table S2); thinnest thickness for effective EA in the Ku and X band (Table S3). See DOI: 10.1039/c8ra05843a



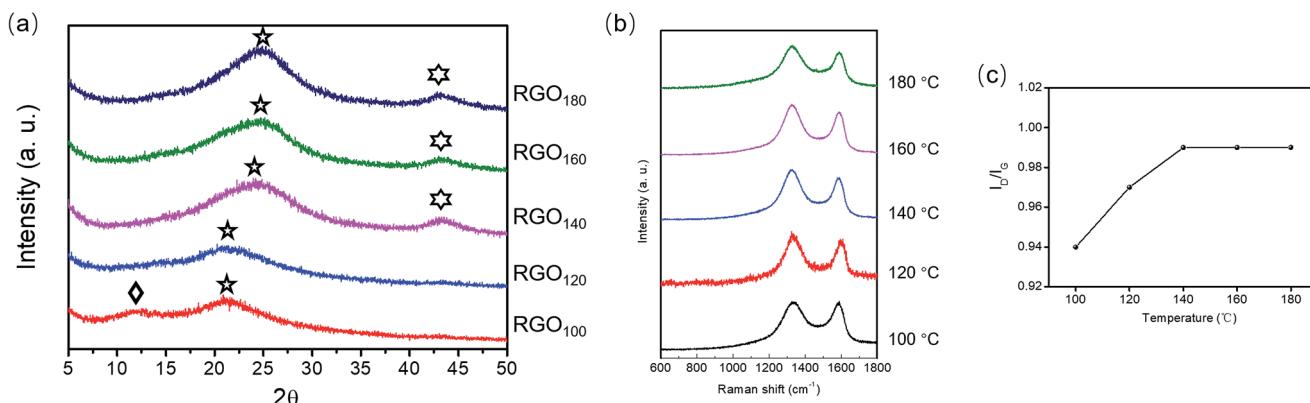


Fig. 1 XRD (a), Raman (b) and I_D/I_G (c) spectra of RGO₁₀₀, RGO₁₂₀, RGO₁₄₀, RGO₁₆₀ and RGO₁₈₀.

contrary, RGO₁₄₀, RGO₁₆₀ and RGO₁₈₀ show the diffraction peaks of graphite (100) around 42°, suggesting that a higher degree of graphitization can be achieved by increasing the hydrothermal temperature. A typical and strong D and G band around 1347 and 1581 cm⁻¹ are shown in the Raman spectra of the as-prepared samples (Fig. 1b). On the one hand, removal of oxygen containing groups in GO makes the G band separate from the D band;¹⁶ on the other hand, defects are produced in the lattice structure during the removal of oxygen containing groups, because of the increase of I_D/I_G (Fig. 1c).⁹

The surface chemistry of RGO₁₀₀, RGO₁₂₀, RGO₁₄₀, RGO₁₆₀ and RGO₁₈₀ was measured by XPS. The C/O atomic ratios increase from 6.35 to 9.44 (Table S1† and Fig. 2f), which suggests that the degree of GO reduction has a positive correlation with the hydrothermal temperature. The C 1s spectra of each RGO sample are shown in Fig. 2, and three typical carbon bonds can be found in each sample, including C-C (~284.8 eV), C-O

(~287.0 eV), and O=C-O (~288.5 eV).¹ Signals resulting from the decrease of C-O are more significant than those of O=C-O, which suggests that most of the oxygen functional groups are removed from the surface of the GO rather than the edges.^{17,18}

Fig. 3 shows the curves of complex permittivity (ϵ_r) of the as-prepared RGO₁₀₀, RGO₁₂₀, RGO₁₄₀, RGO₁₆₀ and RGO₁₈₀/wax composites, where the filler loading ratios of each RGO sample are 3, 5 and 7 wt%, respectively. With the increase to the loading ratio of each RGO sample, both the ϵ' and ϵ'' values increase significantly, which means that increasing the loading ratio can enhance the dielectric properties of the RGO/wax composites. This phenomenon can be well explained according to the effective medium theory.⁷ Meanwhile, the values of ϵ' and ϵ'' under the same filler loading ratios also increase gradually with the increase of hydrothermal temperature. It reveals that tunable permittivity of the composites can be achieved through controlling the reaction temperature of GO.

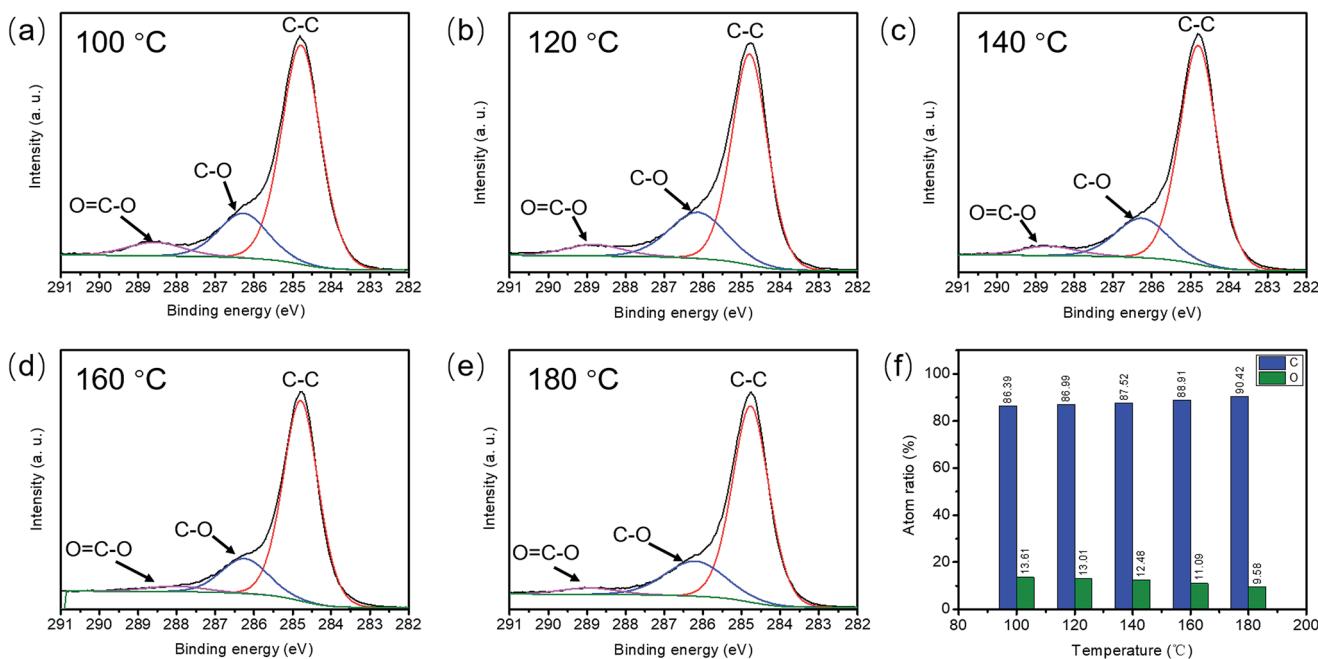


Fig. 2 C 1s spectra of RGO₁₀₀ (a), RGO₁₂₀ (b), RGO₁₄₀ (c), RGO₁₆₀ (d) and RGO₁₈₀ (e) using XPS, and the ratios of C/O in each sample (f).

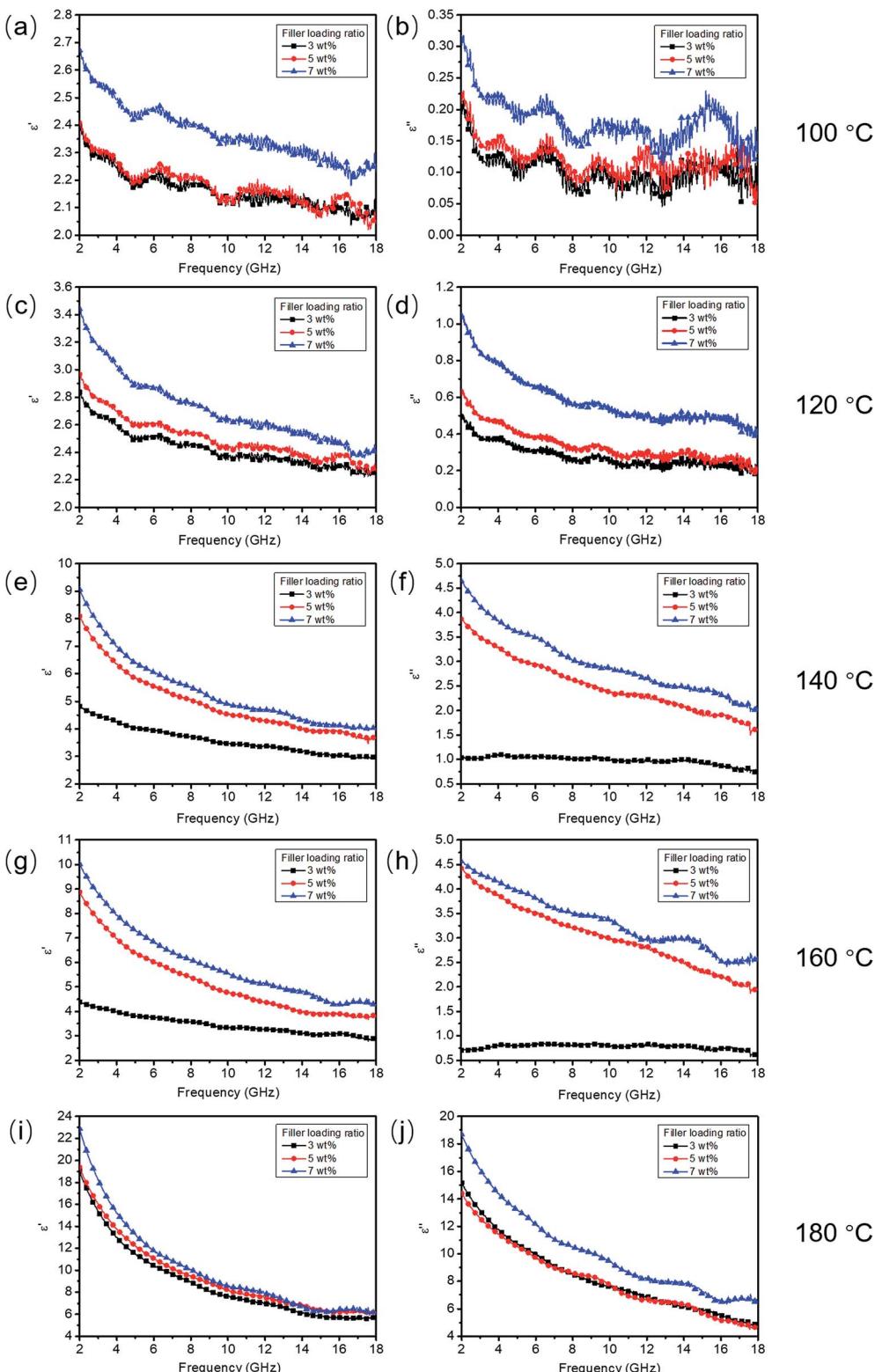


Fig. 3 Frequency-dependent real and imaginary permittivity of RGO_{100} (a and b), RGO_{120} (c and d), RGO_{140} (e and f), RGO_{160} (g and h), and RGO_{180} (i and j), with filler loading ratios of 3, 5 and 7 wt% in the wax composites.

The optimal thicknesses of each sample were taken into account for higher EA performances, and the results are shown in Fig. 4. Under the same loading ratio of RGO, the EA performance of each sample becomes more excellent as the

hydrothermal temperature of GO increases up until 160 °C. However, when the reaction temperature rises to 180 °C, the EA performances of the composites become worse than before. For each sample, the value for the strongest absorption point (RL_{str})

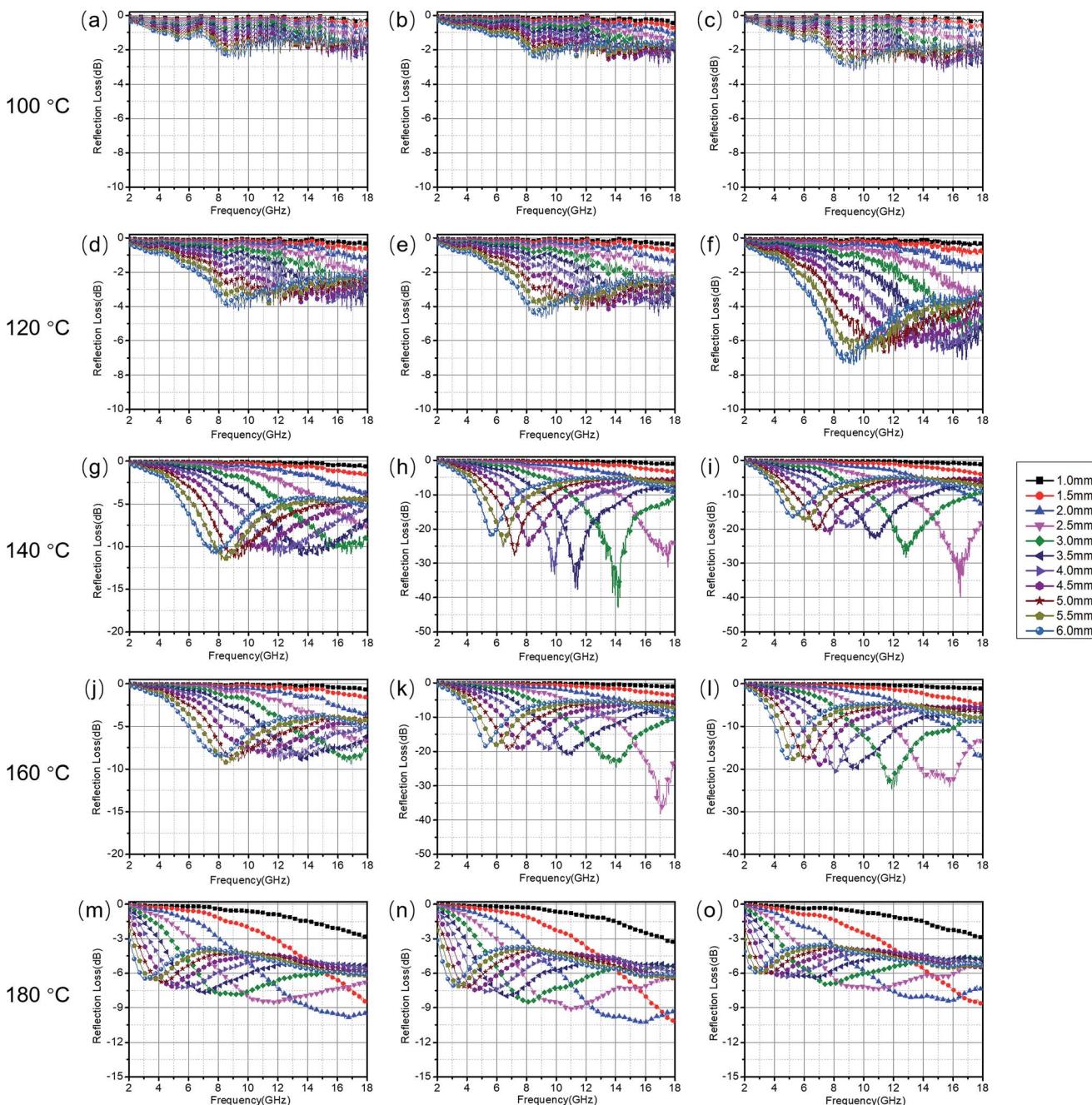


Fig. 4 RL curves of RGO_{100} (a–c), RGO_{120} (d–f), RGO_{140} (g–i), RGO_{160} (j–l) and RGO_{180} (m–o) with filler loading ratios of 3, 5 and 7 wt% in wax composites.

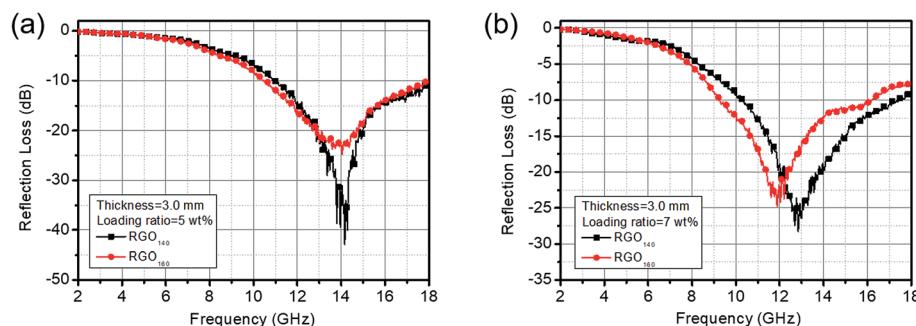


Fig. 5 A comparison of the effective EA bandwidths of composites loaded with 5 wt% and 7 wt% RGO_{140} and RGO_{160} at the same thickness of 3 mm.

and the absorption bandwidth gradually shift toward a lower frequency. This can be explained by the consideration that formation of the quarter-wavelength attenuation requires the absorbing thickness to meet the phase match conditions.⁹ It can be found that the composites loaded with RGO₁₄₀ and

RGO₁₆₀ possess much better EA performances than the other samples. Table S2[†] summarizes the typical thickness dependent EA performance of the RGO₁₄₀ and RGO₁₆₀/wax composites with the filler loading ratios of 5 and 7 wt%. The composites loaded with 5 wt% RGO₁₄₀ and RGO₁₆₀ respectively show effective EA bandwidths (lower than -10 dB) of 7.04 and 7.56 GHz with RL_{str} values of -42.86 and -24.78 dB, both at a thickness of 3 mm (Fig. 5a). When the filler loading ratio increased to 7 wt%, the RGO₁₄₀ and RGO₁₆₀/wax composites show effective EA bandwidths of 7.12 and 7.00 GHz with RL_{str} values of -28.25 and -24.78 dB, respectively, which are also under a thickness of 3 mm (Fig. 5b). From further calculation results, the composite loaded with 5 wt% RGO₁₆₀ has the broadest effective EA bandwidth of 7.56 GHz from 2.9 to 3.2 mm (Fig. 6). When compared with many reported studies (Table S3[†]), it is hard to find a material that can demonstrate such a broadband EA performance at such a low filler loading ratio. The results prove that pure RGO can also demonstrate an ideal electromagnetic absorption performance without any other nanomaterials through a most acceptable method.

The low values of ϵ'' lead to the poor EA performances of the RGO₁₀₀ and RGO₁₂₀/wax composites.¹⁹ The values of ϵ'' are gradually enhanced in the RGO₁₄₀, RGO₁₆₀ and RGO₁₈₀/wax composites, however, the EA performance of the RGO₁₈₀/wax

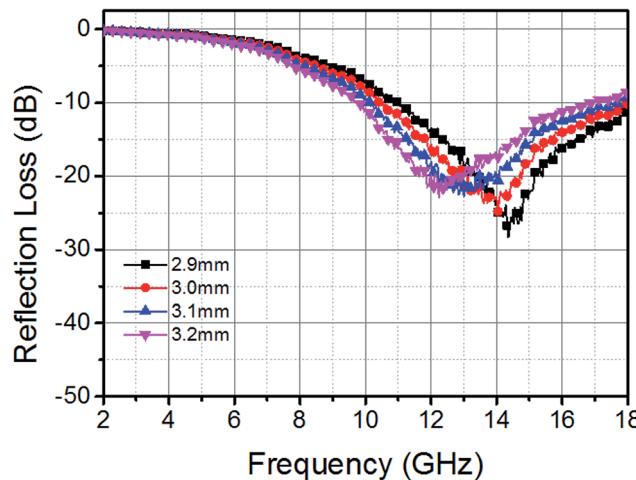


Fig. 6 RL curves of RGO₁₆₀/wax composites, where the filler loading ratio is 5 wt% and the thicknesses are from 2.9 to 3.2 mm.

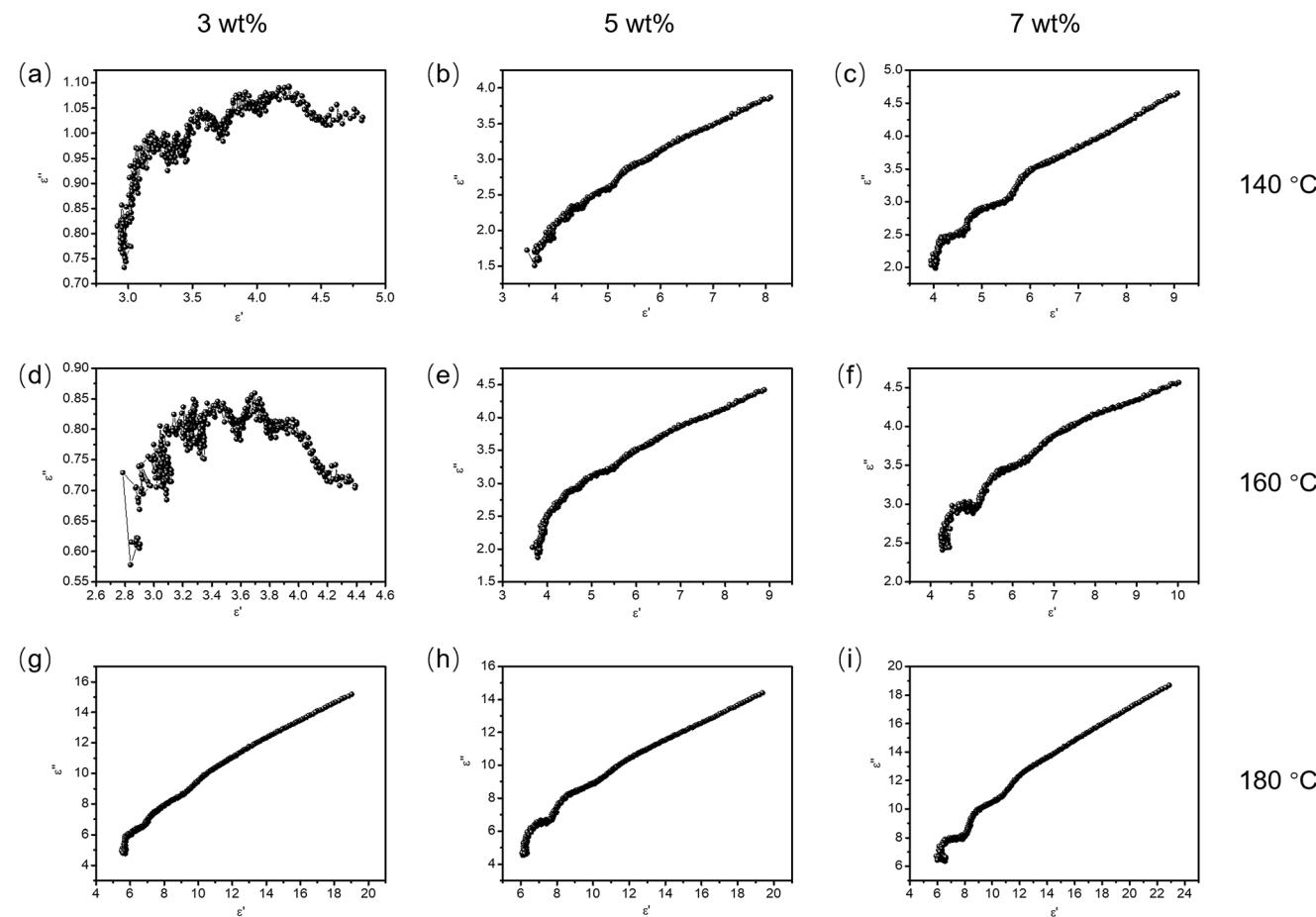


Fig. 7 Cole–Cole plots for the RGO₁₄₀/wax (a–c), RGO₁₆₀/wax (d–f), and RGO₁₈₀/wax (g–i) composites.

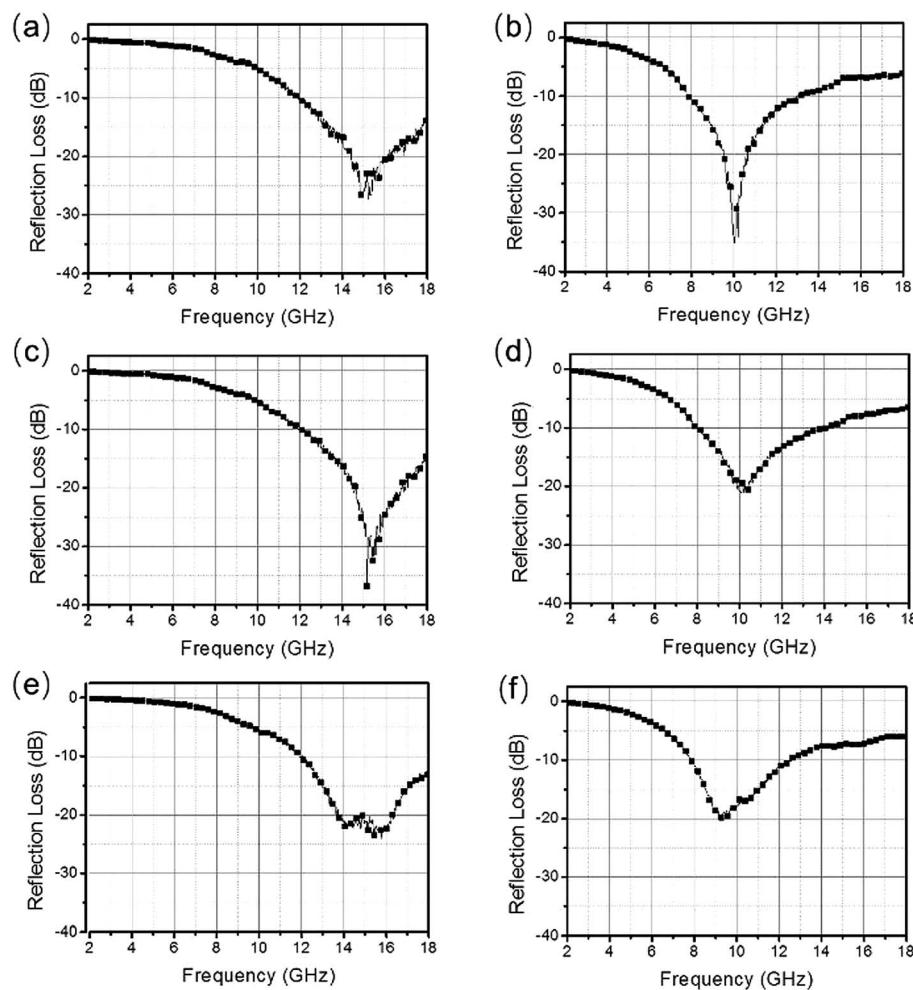


Fig. 8 The thinnest thicknesses to absorb Ku and X bands effectively: RL curves of a composite loaded with 5 wt% RGO_{140} (a and b) and RGO_{160} (c and d); RL curves of a composite loaded with 7 wt% RGO_{160} .

composite is not strengthened further. In the Cole–Cole curves (Fig. 7) of the RGO_{140} and RGO_{160} /wax composites, several semicircle-like structures can be obviously found in the composites with a filler loading ratio of 3 wt%. On the contrary, no evident semicircle is found in the Cole–Cole curve of the RGO_{180} /wax composite, as well as the RGO_{140} and RGO_{160} /wax composites with filler loading ratios of 5 and 7 wt%. Instead, it has the tendency to become linear. This reveals that conductivity loss plays the main role in the dielectric loss of this composite, and polarization induced by defects, chemical bonds and interfaces is hidden by the conductivity loss.⁹ Increased conductivity can provide more efficient EA in theory, however, electromagnetic waves may not be able to enter the composite, due to the impedance of the composite which is far less than that of free space.²⁰ A composite has a much higher dielectric loss than others under a very low filler loading ratio of RGO_{180} , thus the impedance of the composite will be further depressed with the increased filler loading ratio. Finally, the well-reduced RGO_{180} loses its advantage for EA.

In practice, as an ideal EA material, not only is a broad and strong absorption bandwidth needed, but an accurate coverage

of the radar channels is also necessary, for example, the Ku band (12.00–18.00 GHz) for satellite broadcasting, and the X band (8.00–12.00 GHz) for synthetic aperture radar and electron spin resonance apparatus.¹ As shown in Fig. 8 and Table S4,[†] both RGO_{140} and RGO_{160} composites with loading ratios of 5 wt% can effectively dissipate incident electromagnetic energy in the whole Ku band at thicknesses of 2.8 and 2.7 mm, as well as the X band at thicknesses of 3.9 and 3.7 mm. Besides 5 wt% loaded RGO, when RGO_{160} is loaded at 7 wt% in the composite, thinner thicknesses of 2.5 and 3.5 mm are required for effective EA in the Ku and X bands, respectively.

Conclusions

In summary, we have studied the effects of hydrothermal temperature on the permittivity and EA performance of RGO. With the increase of hydrothermal temperature, oxygen functional groups can be removed from GO effectively, especially those from the edges of GO. The composites loaded with 5 wt% RGO_{140} and RGO_{160} respectively show effective EA bandwidths of 7.04 and 7.56 GHz with RL_{str} values of -42.86 and -24.78 dB, both at

thicknesses of 3 mm. When the filler loading ratio is increased to 7 wt%, the RGO_{140} and RGO_{160} /wax composites respectively show effective EA bandwidths of 7.12 and 7.00 GHz with RL_{str} values of -28.25 and -24.78 dB, also at thicknesses of 3 mm.

Statement of contributions

Fan Wu, Aming Xie and Zhangqi Feng designed the experiments. Yilu Xia, Jiankun Wang and Chaochan Chen carried out the synthetic experiments. Da Huo, Yue Wen and Wenyue Wang carried out the characterization. Mengxiao Sun and Chang Xu calculated the RL curves. Yilu Xia, Jiankun Wang and Chaochan Chen wrote this paper.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was financially supported by the National Natural Science Foundation of China (51403236, 51702161), the Fundamental Research Funds for the Central University (AE89926) and the Natural Science Foundation of Jiangsu Province (BK20161466).

Notes and references

- 1 F. Wu, Q. Zeng, Y. Xia, M. Sun and A. Xie, *Appl. Phys. Lett.*, 2018, **112**, 192902.
- 2 F. Meng, H. Wang, F. Huang, Y. Guo, Z. Wang, D. Hui and Z. Zhou, *Composites, Part B*, 2018, **137**, 260.
- 3 B. Quan, X. Liang, G. Ji, Y. Cheng, W. Liu, J. Ma, Y. Zhang, D. Li and G. Xu, *J. Alloys Compd.*, 2017, **728**, 1065.

- 4 W. Cao, X. Wang, J. Yuan, W. Wang and M. Cao, *J. Mater. Chem. C*, 2015, **3**, 10017.
- 5 B. Wu, H. Tuncer, A. Katsounaros, W. Wu, M. Cole, K. Ying, L. Zhang, W. I. Milne and Y. Hao, *Carbon*, 2014, **77**, 814.
- 6 C. Chen, N. Pu, Y. Liu, S. Huang, C. Wu, M. Ger, Y. Gong and Y. Chou, *Composites, Part B*, 2017, **114**, 395.
- 7 M. Ning, M. Lu, J. Li, Z. Chen, Y. Dou, C. Wang, F. Rehman, M. Cao and H. Jin, *Nanoscale*, 2015, **7**, 15734.
- 8 F. Wu, A. Xie, M. Sun, W. Jiang and K. Zhang, *Mater. Lett.*, 2017, **193**, 30.
- 9 F. Wu, A. Xie, M. Sun, Y. Wang and M. Wang, *J. Mater. Chem. A*, 2015, **3**, 14358.
- 10 F. Wu, Y. L. Xia, Y. Wang and M. Y. Wang, *J. Mater. Chem. A*, 2014, **2**, 20307.
- 11 Y. Wang, D. L. Chen, X. Yin, P. Xu, F. Wu and M. He, *ACS Appl. Mater. Interfaces*, 2015, **7**, 26226.
- 12 Y. Zhang, X. Wang and M. Cao, *Nano Res.*, 2018, **11**, 1426.
- 13 M. Li, X. Cao, S. Zheng and S. Qi, *J. Mater. Sci.*, 2017, **28**, 16802.
- 14 S. Zhou, Y. Huang, J. Yan, X. Han and X. Chen, *J. Mater. Sci.*, 2017, **28**, 18558.
- 15 K. Palanisamy, J. H. Um, M. Jeong and W. Yoon, *Sci. Rep.*, 2016, **6**, 31275.
- 16 A. C. Ferrari and D. M. Basko, *Nat. Nanotechnol.*, 2013, **8**, 235.
- 17 A. Lierf, H. Y. He, M. Forster and J. Klinowski, *J. Phys. Chem. B*, 1998, **102**, 4477.
- 18 H. Y. He, J. Klinowski, M. Forster and A. Lierf, *Chem. Phys. Lett.*, 1998, **287**, 53.
- 19 X. Yin, L. Kong, L. Zhang, L. Cheng, N. Travitzky and P. Greil, *Int. Mater. Rev.*, 2014, **59**, 326.
- 20 A. Xie, F. Wu, M. Sun, X. Dai, Z. Xu, Y. Qiu, Y. Wang and M. Wang, *Appl. Phys. Lett.*, 2015, **106**, 222902.

